CLAIM AMENDMENTS

- 1. (Currently Amended) A method of identifying a nucleic acid in a sample, comprising:
 - a) combining the sample with a polynucleotide probe comprising a sequence identical or complementary to at least 10 consecutive nucleotides contained in SEQ ID NO:224, such that the probe hybridizes specifically to the nucleic acid if the nucleic acid encodes human telomerase reverse transcriptase (hTRT) or fragment thereof;
 - b) detecting any hybrid formed as a result of a); and
 - c) identifying the nucleic acid as encoding at least a portion of human telemerase reverse transcriptase (hTRT) hTRT or fragment thereof if the hybrid is detected;

wherein the probe hybridizes specifically to a DNA having the sequence of the hTRT encoding region of SEO, ID NO:224 at 5°C to 25°C below T_m in aqueous solution at 1 M NaCl;

wherein T_m is the melting temperature of double-stranded DNA having the sequence of said encoding region under the same reaction conditions.

- (Currently Amended) A method of detecting a nucleic acid that encodes hTRT or fragment thereof in a sample, comprising:
 - a) combining the sample with a polynucleotide probe such that the probe hybridizes specifically to a nucleic acid comprising at least 100 consecutive nucleotides contained in SEQ ID NO:224 if present in the sample; and
 - b) detecting any hybrid formed as a result of a);

wherein the polynucleotide probe comprises consists essentially of a sequence identical or complementary to at least 25 or more consecutive nucleotides contained in from the hTRT encoding region of SEQ ID NO:224.

- 3. (Original) The method of claim 2, wherein the hTRT nucleic acid is human genomic DNA.
- (Currently Amended) The method of claim 2, wherein the hTRT nucleic acid is human-mRNA mRNA or cDNA.
- (Currently Amended) The method of claim 2, wherein the hTRT nucleic acid comprises at least 250 or more nucleotides of SEQ ID NO:224.
- (Currently Amended) The method of claim 2, wherein the hTRT nucleic acid comprises at least 500 or more nucleotides of SEQ ID NO:224.

- 7. (Currently Amended) The method of claim 2, wherein the probe comprises a sequence identical or complementary to at least-30 or more consecutive nucleotides contained in from the hTRT encoding region of SEQ ID NO:224.
- 8. (Currently Amended) The method of claim 2, wherein the probe comprises a sequence identical or complementary to at-least 50 or more consecutive nucleotides contained in from the hTRT encoding region of SEQ ID NO:224.
- (Currently Amended) The method of claim 2, wherein the probe comprises a sequence identical
 or complementary to at least 100 or more consecutive nucleotides centained in from the hTRT
 encoding region of SEQ ID NO:224.
- 10. (Original) The method of claim 2, wherein the probe comprises a sequence not contained in SEQ. ID NO:62.
- 11. (Original) The method of claim 9, wherein the probe comprises a sequence not contained in SEQ. ID NO:62.
- 12. (Original) The method of claim 2, wherein the sample is a human biological sample.
- 13. (Currently Amended) A method of identifying a nucleic acid in a sample, comprising:
 - a) combining the sample with a polynucleotide primer containing a sequence identical or complementary to at least 10 consecutive nucleotides contained in SEQ ID NO:224, under conditions that the primer amplifies specifically primes amplification of the nucleic acid if the nucleic acid encodes human telemerase reverse transcriptase (hTRT) or fragment thereof;
 - b) detecting any amplification product formed as a result of a); and
 - c) identifying the nucleic acid as encoding at least a portion of hTRT or fragment thereof if the amplification product is detected;

wherein the primer hybridizes specifically to a DNA having the sequence of the hTRT encoding region of SEQ. ID NO:224 at 5°C to 25°C below T_m in aqueous solution at 1 M NaCl:

wherein T_m is the melting temperature of double-stranded DNA having the sequence of said encoding region under the same reaction conditions.

- 14. (Currently Amended) A method of detecting a nucleic acid encoding at least a portion of hTRT or fragment thereof in a sample, comprising:
 - a) combining the sample with a polynucleotide primer such that the primer amplifice polynucleotide primers so as to prime amplification of nucleic acid encoding at least a portion of hTRT or fragment thereof if present in the sample; and
 - b) detecting any amplified product formed as a result of a);

wherein the polynucleotide primer_comprises each of said primers consists essentially of a sequence identical or complementary to at-least 16 or more consecutive nucleotides contained in from the hTRT encoding region of SEQ ID NO:224.

- 15. (Currently Amended) The method of claim 14, wherein the polynucleotide-primer comprises each of said primers consists essentially of a sequence identical or complementary to at least 30 or more consecutive nucleotides contained in from the hTRT encoding region of SEQ ID NO:224.
- 16. (Currently Amended) The method of claim 14, wherein the polynucleotide primer comprises each of said primers consists essentially of a sequence identical or complementary to at least 50 or more consecutive nucleotides centained in from the hTRT encoding region of SEQ ID NO:224.
- 17. (Original) The method of claim 14, wherein the sample is a human biological sample.
- 18. (Original) The method of claim 14, wherein the sample comprises human genomic DNA.
- (Currently Amended) The method of claim 14, wherein the sample comprises human mRNA hTRT mRNA or cDNA.
- 20. CANCELLED.
- (Original) The method of claim 14, wherein the primer-comprises primers comprise a sequence not contained in SEQ. ID NO:62.
- 22. CANCELLED

- 23. (Withdrawn) (Currently Amended) A combination of oligonucleotide primers for PCR amplification for use in detecting an hTRT nucleic acid according to claim 14, comprising a first primer that hybridizes to a polynucleotide consisting of SEQ ID NO:224 under stringent amplification conditione, and a second primer that hybridizes to the complement of said nucleic acid under stringent amplification conditions wherein each primer consists essentially of a sequence identical or complementary to 15 or more consecutive nucleotides from the hTRT encoding region of SEQ ID NO:224.
- 24. (Withdrawn) The combination of primers of claim 23, wherein either each primer comprises between consists of 15-30 nucleotides.
- (Withdrawn) The combination of primers of claim 23, wherein either each primer comprises between consists of 20-25 nucleotides.
- 26. (Withdrawn) The combination of primers of claim 23, wherein 50% or more of the nucleotides of either each primer are guanine and/or cytosine.
- 27. (Withdrawn) (Currently Amended) A PCR product that hybridizes under stringent conditions to a polynucleotide having a sequence consisting of formed while undertaking the detection method of claim 14, comprising 15 or more contiguous nucleotides of the hTRT encoding region of SEQ ID NO:224 or its complement.
- 28. (Withdrawn) (Currently Amended) A hybridization complex formed while undertaking the detection method of claim 2, comprising:
 - a) one strand of a cellular hTRT nucleic acid; and
 - b) one strand of a nucleic acid comprising a recombinant or synthetic fragment of hTEPT; wherein-said fragment of hTRT comprises at least 10 contiguous nucleotides of consisting essentially of 25 or more consecutive nucleotides of the hTRT encoding region of SEQ ID NO:224 or its complement.
- (Withdrawn) The hybridization complex of claim 28, wherein the hTRT nucleic acid is an hTRT mRNA.
- 30. (Withdrawn) The hybridization complex of claim 28, wherein the hTRT nucleic acid is an hTRT cDNA.
- (Withdrawn) The hybridization complex of claim 28, wherein the fragment comprises at least 20 centiguous or more consecutive nucleotides of SEQ ID NO:224 or its complement.

- (Withdrawn) (Currently Amended) The hybridization complex of claim 28, wherein the fragment comprises 10-100 contiguous 100 or more consecutive nucleotides of SEQ ID NO:224 or its complement.
- (Withdrawn) The hybridization complex of claim 28, wherein said hybridization complex is a DNA:DNA complex.
- 34. (Withdrawn) The hybridization complex of claim 28, wherein said hybridization complex is a DNA:RNA complex.
- 35. (New) The method of claim 1, wherein a) comprises combining the sample with the probe at 5°C to 25°C below T_m in aqueous solution at 1 M NaCl.
- 36. (New) The method of claim 1, wherein the hTRT nucleic acid is hTRT mRNA or cDNA.
- 37. (New) The method of claim 1, wherein the probe comprises a sequence identical or complementary to 100 or more consecutive nucleotides from the hTRT encoding region of SEQ ID NO:224.
- 38. (New) The method of claim 1, wherein the probe comprises a sequence not contained in SEQ. ID NO:62.
- 39. (New) The method of claim 1, wherein the sample has been taken from a patient, and the method further comprises determining or assessing a tumor in the patient according to whether a nucleic acid encoding hTRT or an hTRT fragment is detected.
- 40. (New) The method of claim 2, wherein the sample has been taken from a patient, and the method further comprises determining or assessing a tumor in the patient according to whether said nucleic acid hybrid is detected.
- 41. (New) The method of claim 13, wherein a) comprises combining the sample with the primer at 5°C to 25°C below T_m in aqueous solution at 1 M NaCl.
- 42. (New) The method of claim 13, wherein the hTRT nucleic acid is mRNA or cDNA.

- 43. (New) The method of claim 13, wherein the primer comprises a sequence identical or complementary to 30 or more consecutive nucleotides from the hTRT encoding region of SEQ ID NO:224.
- 44. (New) The method of claim 13, wherein the primer comprises a sequence not contained in SEQ. ID NO:62.
- 45. (New) The method of claim 13, wherein the sample has been taken from a patient, and the method comprises determining or assessing a tumor in the patient according to whether a nucleic acid encoding hTRT or an hTRT fragment is detected.
- 46. (New) The method of claim 14, wherein the sample has been taken from a patient, and the method comprises determining or assessing a tumor in the patient according to whether said amplification product is formed.